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IN THE CLAIMS

1-10. (Cancelled)

11. (Currently Amended) Process for the preparation of the amorphous form of methyl (S)-(+)-(2-chlorophenyl)-2-(6, 7-dihydro-4*H*-thieno [3,2- c] pyridine-5-yl-acetate hydrogen-sulfate of the formula

which comprises,

dissolving clopidogrel base in an "A" type a first solvent,

adding

sulfuric acid or

a mixture of sulfuric acid and an "A" or "B" type the first solvent or a second solvent to the mixture,

adding the obtained mixture containing clopidogrel hydrogensulfate to a "B" type the second solvent to obtain a precipitate, and

filtering, optionally washing and drying the obtained precipitate,

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wherein the first solvent is selected from at least one of the group consisting of: an

aprotic solvent that is less polar than the second solvent and a dipolar aprotic solvent, and

wherein the second solvent is selected from at least one of the group consisting of: an

aprotic solvent, a dipolar aprotic solvent and an apolar solvent.

12. (Currently Amended) Process The process according to Claim 1 which comprises

using less polar aprotic solvents preferably claim 11,

wherein in said first solvent,

said aprotic solvent is a halogenated solvents solvent, more preferably dichloromethane,

or and

said dipolar aprotic solvents preferably ketones more preferably lower alkyl ketones.

most preferably acetone, as "A" type solvent solvent is a ketone, and

wherein in said second solvent,

<u>said</u> aprotic solvents preferably solvent is an ether type solvents, more preferably diethyl

ether, tetrahydrofurane, diisopropyl ether, most preferably diisopropyl ether, or,

said dipolar aprotic solvents solvent, preferably is an ester type solvent, more preferably

ethyl acetate, or and

said apolar solvents preferably solvent is an alkyl hydrocarbons more preferably

cyclohexane, hexane, heptane, most preferably cyclohexane as "B" type solvent hydrocarbon.

13. (Currently Amended) Process The process according to Claim 1-which comprises claim

11, wherein the method comprises:

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dissolving [[of]] clopidogrel base in dichloromethane to make a solution, adding sulfuric acid to the solution, mixing the obtained solution with cyclohexane to form a precipitate, then and filtering the obtained precipitate.

- 14. (New) The process according to claim 11, which further comprises: washing the precipitate; and drying the precipitate.
 - 15. (New) The process according to claim 11,
 wherein in said first solvent,
 said aprotic solvent is dichloromethane, and
 said dipolar aprotic solvent is a lower alkyl ketone, and
 wherein in said second solvent,

said aprotic solvent is selected from the group consisting of: diethyl ether, tetrahydrofuran and diisopropyl ether,

said dipolar aprotic solvent is ethyl acetate, and

said apolar solvent is selected from the group consisting of cyclohexane, hexane and heptane.

16. (New) The process according to claim 11, wherein in said first solvent,

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said aprotic solvent is dichloromethane, and

said dipolar aprotic solvent is acetone, and

wherein in said second solvent,

said aprotic solvent is diisopropyl ether,

said dipolar aprotic solvent is ethyl acetate, and

said apolar solvent is cyclohexane.

17. (New) The process according to claim 11, wherein the first solvent is acetone and

wherein the second solvent is diisopropyl ether.

18. (New) The process according to claim 11, wherein the first solvent is

dichloromethane and wherein the second solvent is diisopropyl ether.

19. (New) The process according to claim 11, wherein the first solvent is

dichloromethane and wherein the second solvent is ethyl acetate.

20. (New) The process according to claim 11, wherein said first solvent is present in an

amount not greater than 37 ml per gram of clopidogrel base.

21. (New) The process according to claim 11, wherein said first solvent is present in an

amount of between 31 and 37 ml per gram of clopidogrel base.